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RNA, or heterogeneous nuclear ribonucleoprotein A2/B1 RNA.in an animal, the method comprising the steps of detecting an amplified product according to claim 2 and detecting or diagnosing a disease associated with epidermal growth factor RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, or heterogeneous nuclear ribonucleoprotein A2/B1 RNA.

- 19. The method of claim 17 wherein the disease is a malignancy or premalignancy.
- 20. The method of claim 18 wherein the disease is a malignancy or premalignancy.
- A method for monitoring an animal or human for a malignant or premalignant disease, wherein the malignant or premalignant disease is associated with a tumor-derived or tumor-associated RNA that is epidermal growth factor RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, or heterogeneous nuclear ribonucleoprotein A2/B1 RNA, or any combination thereof, the method comprising the step of:
 - qualitatively or quantitatively, wherein the RNA is epidermal growth factor
 RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, or
 heterogeneous nuclear ribonucleoprotein A2/B1 RNA or any combination thereof,
 according to a method comprising the steps of:
 - a) extracting mammalian RNA from plasma or serum, wherein a fraction of said extracted RNA comprises epidermal growth factor RNA, epidermal growth factor

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- receptor RNA, her-2/neu RNA, c-myc RNA, heterogeneous nuclear ribonucleoprotein A2/B1 RNA or any combination thereof;
- b) amplifying or signal amplifying said fraction of the extracted RNA or corresponding cDNA, wherein amplification is performed qualitatively or quantitatively using primers specific for the tumor-derived or tumor-associated RNA or cDNA corresponding thereto, to produce an amplified product; and
- c) detecting the amplified product produced from RNA or cDNA corresponding thereto.
- 22. A method for monitoring an animal or human for a malignant or premalignant disease, wherein the malignant or premalignant disease is associated with tumor-derived or tumor-associated RNA that is epidermal growth factor RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, or heterogeneous nuclear ribonucleoprotein A2/B1 RNA or any combination thereof, the method comprising the step of:
 - detecting qualitatively or quantitatively RNA associated with the malignant or premalignant disease, wherein the RNA is epidermal growth factor RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, or heterogeneous nuclear ribonucleoprotein A2/B1 RNA or any combination thereof, according to a method comprising the steps of:
 - a) extracting mammalian RNA from a bodily fluid, wherein a fraction of said extracted RNA comprises epidermal growth factor RNA, epidermal growth factor receptor RNA, her-2/neu RNA, c-myc RNA, heterogeneous nuclear ribonucleoprotein A2/B1 RNA or any combination thereof;

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- b) amplifying or signal amplifying said fraction of the extracted RNA or cDNA corresponding thereto, wherein amplification is performed qualitatively or quantitatively using primers specific for the tumor-derived or tumor-associated RNA or cDNA corresponding thereto, to produce an amplified product; and
- c) detecting the amplified product produced from said RNA or cDNA corresponding thereto.
- 23. A method for selecting an animal or human with cancer for an epidermal growth factor-directed therapy comprising the step of performing the method of claim 1 using blood plasma or serum from said animal or human and selecting the animal or human for an epidermal growth factor-directed therapy when epidermal growth factor RNA is detected in the animal or human's plasma or serum.
- 24. A method for selecting an animal or human with cancer for an epidermal growth factor-directed therapy comprising the step of performing the method of claim 2 using a bodily fluid from said animal or human and selecting the animal or human for an epidermal growth factor-directed therapy when epidermal growth factor RNA is detected in the animal or human's plasma or serum.
- 20 25. A method for selecting an animal or human with cancer for an epidermal growth factor receptor-directed therapy comprising the step of performing the method of claim 1 using blood plasma or serum from said animal or human and selecting the animal or human for